PATENTS

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

APPLICANTS: CHOI ET AL-1 (PCT)

PCT NO.: PCT/KR2005/002170 PCT FILED: JULY 6, 2005

SERIAL NO.: 10/589,328

TITLE: DIAGNOSIS METHOD AND KITS FOR INHERITED NEUROPATHIES CAUSED BY DUPLICATION OR DELETION OF CHROMOSOME 17P11.2-P12 REGION

SUPPLEMENTAL INFORMATION DISCLOSURE STATEMENT

MAIL STOP PCT
Commissioner of Patents
P.O. Box 1450
Alexandria, VA 22313-1450

Dear Sir:

Supplemental to the Information Disclosure Statement filed August 14, 2006, Applicants wish to submit to the Patent Examiner copies of six references cited in the Specification and listed on the previously filed PTO Form-1449, but not previously submitted. A new PTO Form 1449 listing these references is enclosed.

Because this Information Disclosure Statement (IDS) is being filed before the issuance of a first Office Action on the merits, it is believed that no fee is due. However, the Commissioner is hereby authorized to charge Deposit Account No. 03-2468 for any additional fees or credit any overpayment in connection with this IDS. It is respectfully requested that the foregoing IDS be incorporated into the official file of the present patent application.

Respectfully submitted,
Byung ~~Ok~~ CHOI ET AL.

COLLARD & ROE, P.C.
1077 Northern Boulevard
Roslyn, New York 11576
(516) 365-9802

Frederick J. Dorchak, Reg.No. 29,298
Edward R. Freedman, Reg.No. 26,048
Attorneys for Applicant

FJD:cmm
Enclosures: PTO Form-1449 and 6 references

I hereby certify that this correspondence is being deposited with the U.S. Postal Service as first class mail in an envelope addressed to: Commissioner of Patents, P.O. Box 1450, Alexandria, VA 22313-1450, on November 20, 2006.

Kelly Espitia

FORM PTO-1449 (REV. 7-80) | U.S. DEPARTMENT OF COMMERCE PATENT AND TRADEMARK OFFICE

ATTY. DOCKET NO. : CHOI ET AL--1 PCT

SERIAL NO. 10/589,328

LIST OF REFERENCES CITED BY APPLICANT

(Use several sheets if necessary)

APPLICANT : Byung Ok CHOI ET AL.

FILING DATE:

GROUP:

U.S. PATENT DOCUMENTS

*EXAMINER INITIAL		DOCUMENT NUMBER	DATE	NAME	CLASS	SUBCLASS	FILING DATE IF APPROPRIATE
	AA						
	AB						
	AC						
	AD						

OTHER REFERENCES *(Including Author, Title, Date, Pertinent Pages, Etc.)*

	AE		Berger P., Young P., Suter U. "Molecular cell biology of Charcot-Marie-Tooth disease," Neurogenet. 4: 1-15 (2002).
	AF		Sereda M.W., Horste G.M., Suter U., Uzma N., Nave K.-A. "Therapeutic administration of progesterone antagonist in a model of Charcot-Marie-Tooth disease," Nature Genet. 9: 1533-1537 (2003).
	AG		Passage E., Norreel J.C., Noach-Fraissignes P., Sanguedolce V., Pizant J., Thirion X., Robaglia-Schlupp A., Pellissier J.F., Fontes M. "Ascorbic acid treatment corrects the phenotype of a mouse model of Charcot-Marie-Tooth disease," Nature Genet. 10: 396-401 (2004).
	AH		Mersiyanova I.V., Ismailov S.M., Polyakov A.V., Dadali E.L., Fedotov V.P., Nelis E., et al. "Screening for Mutations in the Peripheral Myelin Genes PMP22, MPZ and Cx32 (GJB1) in Russian Charcot-Marie-Tooth Neuropathy Patients," Human Mutat. 15: 340-347 (2000).
	AI		Yoshihara T., Yamamoto M., Doyu M., Misu K.I., Hattori N. Hasegawa Y., Mokuno K., Mitsuma T., Sobue G. "Mutations in the Peripheral Myelin Protein Zero and Connexin32 Genes Detected by non-Isotopic RNase Cleavage Assay and Their Phenotypes in Japanese Patients with Charcot-Marie-Tooth Disease," Hum. Mutat. 16: 177-178 (2000).
	AJ		Numakura C., Lin C., Ikegami T., Guldberg P., Hayasaka K. "Molecular Analysis in Japanese Patients With Charcot-Marie-Tooth Disease: DGGE Analysis for PMP22, MPZ and Cx32/GJB1 Mutations," Human Mutat. 20: 392-398 (2002).
	AK		
	AL		

EXAMINER | DATE CONSIDERED

*EXAMINER: Initial if reference considered, whether or not citation is in conformance with MPEP 609; draw line through citation if not in conformance and not considered. Include copy of this form with next communication to applicant.